WHAT IS CLAIMED IS:

1. A compound of the formula:

Formula I

wherein:

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R¹ is optionally substituted lower alkyl, optionally substituted cycloalkyl, optionally substituted aryl, or optionally substituted heteroaryl;

X is a covalent bond or optionally substituted alkylene;

10 R^2 is R^4 -Z-Y-C=C- or optionally substituted pyrazolyl:

in which Y is optionally substituted alkylene, Z is oxygen, sulfur or -NH-, and R⁴ is optionally substituted aryl or optionally substituted heteroaryl; and

R³ is hydroxymethyl or -C(O)-NR⁵R⁶;

in which R⁵ and R⁶ are independently hydrogen or lower alkyl.

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- 2. The compound of claim 1, wherein R² is optionally substituted pyrazol-1-yl.
- 3. The compound of claim 2, wherein R^1 is optionally substituted alkyl or optionally substituted aryl and R^3 is hydroxymethyl.

- 4. The compound of claim 3, wherein R² is pyrazo-1-yl substituted by optionally substituted lower alkyl, ester, aminocarbonyl, optionally substituted aryl, or optionally substituted heteroaryl.
- 5. The compound of claim 4, wherein pyrazol-1-yl is substituted by optionally substituted phenyl or optionally substituted benzyl.

- 6. The compound of claim 5, wherein R¹ is optionally substituted lower alkyl and X is a covalent bond.
- 7. The compound of claim 6, wherein R¹ is methyl and R² is 4-(45 methoxyphenyl)pyrazol-1-yl, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-{2-[4-(4methoxyphenyl)pyrazolyl]-6-(methylamino)purin-9-yl}oxolane-3,4-diol.
 - 8. The compound of claim 6, wherein R¹ is n-propyl and R² is 4-(4-methoxyphenyl)pyrazol-1-yl, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-{2-[4-(4-methoxyphenyl)pyrazolyl]-6-(n-propylamino)purin-9-yl}oxolane-3,4-diol.
 - 9. The compound of claim 6, wherein R¹ is methyl and R² is 4-(4-chlorobenzylaminocarbonyl)pyrazol-1-yl, namely (1-{9-[(4S,2R,3R,5R)-3,4-dihydroxy-5-(hydroxymethyl)oxolan-2-yl]-6-(methylamino)purin-2-yl}pyrazol-4-yl)-N-(4-chlorophenyl)carboxamide.
 - 10. The compound of claim 6, wherein R¹ is methyl and R² is 4-(4-chlorobenzylaminocarbonyl)pyrazol-1-yl, namely (1-{9-[(4S,2R,3R,5R)-3,4-dihydroxy-5-(hydroxymethyl)oxolan-2-yl]-6-(methylamino)purin-2-yl}pyrazol-4-yl)-N-(4-chlorophenyl)carboxamide.
 - 11. The compound of claim 4, wherein R² is pyrazo-1-yl substituted by optionally substituted heteroaryl.
- 25 12. The compound of claim 11, wherein R¹ is n-propyl and R² is 4-(pyrid-2-yl)pyrazol-1-yl, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-[4-(pyridin-2-yl)pyrazolyl]-6-(n-propylamino)purin-9-yl}oxolane-3,4-diol.
 - 13. The compound of claim 5, wherein R¹ is optionally substituted aryl and X is alkylene.

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- 14. The compound of claim 13, wherein R¹ is 3-iodobenzyl and R² is 4-(4-methoxyphenyl)pyrazol-1-yl, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-{2-[4-(4-methoxyphenyl)pyrazolyl]-6-(3-iodobenzylamino)purin-9-yl}oxolane-3,4-diol.
- 5 15. The compound of claim 1, wherein R² is optionally substituted pyrazol-4-yl.
 - 16. The compound of claim 15, wherein R^1 is optionally substituted alkyl or optionally substituted aryl, R^3 is hydroxymethyl, and X is a covalent bond.
- 17. The compound of claim 16, wherein R¹ is methyl, R² is 1-benzylpyrazol-4-yl, R³ is hydroxymethyl, and X is a covalent bond, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-{2-[1-benzylpyrazolyl]-6-(methylamino)purin-9-yl}oxolane-3,4-diol.
- 18. The compound of claim 16, wherein R¹ is n-propyll, R² is 1-benzylpyrazol-4-yl, R³ is hydroxymethyl, and X is a covalent bond, namely (4S,2R,3R,5R)-5-(hydroxymethyl)-2-{2-[1-benzylpyrazolyl]-6-(n-propylamino)purin-9-yl}oxolane-3,4-diol.
 - 19. The compound of claim 1, wherein R^2 is R^4 -Z-Y-C=C-.
- 20. The compound of claim 19, wherein R⁴ is optionally substituted phenyl and Y is alkylene of 1-3 carbon atoms.
 - 21. The compound of claim 20, wherein R⁴ is phenyl optionally substituted by methoxy or chloro, and Y is methylene.
 - 22. The compound of claim 21, wherein R^1 is optionally substituted alkyl, X is a covalent bond, and R^3 is hydroxymethyl.
- 23. The compound of claim 22, wherein R¹ is methyl, R⁴ is phenyl and Z is oxygen,
 namely 2-hydroxymethyl-5-[6-methylamino-2-(3-phenoxypropyn-1-yl)purin-9-yl]tetrahydrofuran-3,4-diol.

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- 24. A method of treating a disease state in a mammal that is alleviable by treatment with a A₃ adenosine receptor agonist, comprising administering to a mammal in need thereof a therapeutically effective dose of a compound of claim 1.
- 5 25. The method of claim 24, wherein the disease state is cancer.
 - 26. The method of claim 24, wherein the disease state is neutropenia.
- 27. A pharmaceutical composition comprising at least one pharmaceutically acceptable excipient and a therapeutically effective amount of a compound of claim 1.
 - 28. A process for the preparation of a compound of Formula I:

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in which R^2 is optionally substituted pyrazol-1-yl; comprising:

contacting a compound of the formula:

with a compound of formula:

$$\mathbb{R}^5$$
 \mathbb{R}^6
 \mathbb{R}^7
 \mathbb{R}^7

- 29. The process of claim 28, wherein the reaction is conducted in an inert solvent chosen from methanol, ethanol, n-propanol, isopropanol, and t-butanol.
- 30. A process for the preparation of a compound of Formula I:

in which R^2 is optionally substituted pyrazol-4-yl; comprising

10 contacting a compound of the formula:

with a compound of the formula:

in the presence of a palladium complex and a copper salt in an inert solvent, and contacting the product with a mild acid.

- 5 31. The process of claim 30, wherein the palladium complex is Pd(PPh₃)₄, the copper salt is CuI, the inert solvent is N,N-dimethylformamide, and the mild acid is ammonium fluoride.
 - 32. A process for the preparation of a compound of claim 1, in which R² is R⁴-Z-Y-C≡C-; comprising:
- 10 contacting in an inert solvent a compound of the formula:

with a compound of the formula:

in the presence of a mild base, a copper salt and a palladium catalyst.

33. The process of claim 32, wherein the inert solvent is N, N-dimethylformamide, the base is triethylamine, the copper salt is copper iodide, and the palladium catalyst is dichlorobis-(triphenylphosphine)palladium(II).